



- ☐ Philips Holding USA Inc.
- ☐ Philips RS North America Holding Corporation.
- ☐ Polymer Technologies, Inc.
- ☐ Polymer Molded Products LLC.

**II. PLAINTIFF(S)**

2. Name of Plaintiff(s):  
Kelvin Kuntz
- 

3. Name of spouse of Plaintiff (if loss of consortium claim is being made):  
N/A
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4. Name and capacity (*i.e.*, executor, administrator, guardian, conservator, etc.) of other Plaintiff, if any:  
N/A
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5. State(s) of residence of Plaintiff(s) (if the Recalled Device user is deceased, residence at the time of death):  
United States Virgin Islands (St. Thomas)
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**III. DESIGNATED FORUM**

6. Identify the forum (United States District Court and Division) in which the Plaintiff would have filed in the absence of direct filing:  
United States District Court for the Virgin Islands
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**IV. USE OF A RECALLED DEVICE**

7. Plaintiff used the following Recalled Device(s):

<input type="checkbox"/> E30 (Emergency Use Authorization)	<input type="checkbox"/> Dorma 500
<input type="checkbox"/> DreamStation ASV	<input checked="" type="checkbox"/> REMstar SE Auto
<input type="checkbox"/> DreamStation ST, AVAPS	<input type="checkbox"/> Trilogy 100
<input type="checkbox"/> SystemOne ASV4	<input type="checkbox"/> Trilogy 200
<input type="checkbox"/> C-Series ASV	<input type="checkbox"/> Garbin Plus, Aeris, LifeVent
<input type="checkbox"/> C-Series S/T and AVAPS	<input type="checkbox"/> A-Series BiPAP Hybrid A30 (not marketed in U.S.)
<input type="checkbox"/> OmniLab Advanced +	<input type="checkbox"/> A-Series BiPAP V30 Auto
<input type="checkbox"/> SystemOne (Q-Series)	<input type="checkbox"/> A-Series BiPAP A40
<input checked="" type="checkbox"/> DreamStation	<input type="checkbox"/> A-Series BiPAP A30
<input type="checkbox"/> DreamStation Go	<input type="checkbox"/> Other Philips Respironics Device; if other, identify the model:
<input type="checkbox"/> Dorma 400	

**V. INJURIES**

8. Plaintiff alleges the following physical injuries as a result of using a Recalled Device together with the attendant symptoms and consequences associated therewith:

- ☐ COPD (new or worsening)
- ☐ Asthma (new or worsening)
- ☐ Pulmonary Fibrosis
- ☒ Other Pulmonary Damage/Inflammatory Response
- ☐ Cancer \_\_\_\_\_ (specify cancer)
- ☐ Kidney Damage
- ☐ Liver Damage

☐ Heart Damage

☐ Death

☒ Other (specify)

Difficulty sleeping, increased headaches, sinus fluid retention, weakness

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## VI. CAUSES OF ACTION/DAMAGES

9. As to Koninklijke Philips N.V., Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

☐ Count I: Negligence

☒ Count II: Strict Liability: Design Defect

☐ Count III: Negligent Design

☒ Count IV: Strict Liability: Failure to Warn

☒ Count V: Negligent Failure to Warn

☐ Count VI: Negligent Recall

☐ Count VII: Battery

☐ Count VIII: Strict Liability: Manufacturing Defect

☐ Count IX: Negligent Manufacturing

☒ Count X: Breach of Express Warranty

☒ Count XI: Breach of the Implied Warranty of Merchantability

☐ Count XII: Breach of the Implied Warranty of Usability

☒ Count XIII: Fraud

☒ Count XIV: Negligent Misrepresentation

- ☐ Count XV: Negligence Per Se
- ☐ Count XVI: Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
- ☐ Count XVII: Unjust Enrichment
- ☐ Count XVIII: Loss of Consortium
- ☐ Count XIX: Survivorship and Wrongful Death
- ☐ Count XX: Medical Monitoring
- ☐ Count XXI: Punitive Damages
- ☐ Count XXII: Other [specify below]

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10. As to Philips North America LLC, Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

- ☐ Count I: Negligence
- ☒ Count II: Strict Liability: Design Defect
- ☐ Count III: Negligent Design
- ☒ Count IV: Strict Liability: Failure to Warn
- ☒ Count V: Negligent Failure to Warn
- ☐ Count VI: Negligent Recall
- ☐ Count VII: Battery
- ☐ Count VIII: Strict Liability: Manufacturing Defect
- ☐ Count IX: Negligent Manufacturing

- ☒ Count X: Breach of Express Warranty
- ☒ Count XI: Breach of the Implied Warranty of Merchantability
- ☐ Count XII: Breach of the Implied Warranty of Usability
- ☒ Count XIII: Fraud
- ☒ Count XIV: Negligent Misrepresentation
- ☐ Count XV: Negligence Per Se
- ☐ Count XVI: Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
- ☐ Count XVII: Unjust Enrichment
- ☐ Count XVIII: Loss of Consortium
- ☐ Count XIX: Survivorship and Wrongful Death
- ☐ Count XX: Medical Monitoring
- ☐ Count XXI: Punitive Damages
- ☐ Count XXII: Other [specify below]

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11. As to Philips RS North America LLC, Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

- ☐ Count I: Negligence
- ☒ Count II: Strict Liability: Design Defect
- ☐ Count III: Negligent Design
- ☒ Count IV: Strict Liability: Failure to Warn

- ☒ Count V: Negligent Failure to Warn
  - ☐ Count VI: Negligent Recall
  - ☐ Count VII: Battery
  - ☐ Count VIII: Strict Liability: Manufacturing Defect
  - ☐ Count IX: Negligent Manufacturing
  - ☒ Count X: Breach of Express Warranty
  - ☒ Count XI: Breach of the Implied Warranty of Merchantability
  - ☐ Count XII: Breach of the Implied Warranty of Usability
  - ☒ Count XIII: Fraud
  - ☒ Count XIV: Negligent Misrepresentation
  - ☐ Count XV: Negligence Per Se
  - ☐ Count XVI: Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
  - ☐ Count XVII: Unjust Enrichment
  - ☐ Count XVIII: Loss of Consortium
  - ☐ Count XIX: Survivorship and Wrongful Death
  - ☐ Count XX: Medical Monitoring
  - ☐ Count XXI: Punitive Damages
  - ☐ Count XXII: Other [specify below]
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12. As to Philips Holding USA Inc., Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

- ☐ Count I: Negligence
- ☐ Count II: Strict Liability: Design Defect
- ☐ Count III: Negligent Design
- ☐ Count IV: Strict Liability: Failure to Warn
- ☐ Count V: Negligent Failure to Warn
- ☐ Count VI: Negligent Recall
- ☐ Count VII: Battery
- ☐ Count VIII: Strict Liability: Manufacturing Defect
- ☐ Count IX: Negligent Manufacturing
- ☐ Count X: Breach of Express Warranty
- ☐ Count XI: Breach of the Implied Warranty of Merchantability
- ☐ Count XII: Breach of the Implied Warranty of Usability
- ☐ Count XIII: Fraud
- ☐ Count XIV: Negligent Misrepresentation
- ☐ Count XV: Negligence Per Se
- ☐ Count XVI: Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
- ☐ Count XVII: Unjust Enrichment
- ☐ Count XVIII: Loss of Consortium
- ☐ Count XIX: Survivorship and Wrongful Death
- ☐ Count XX: Medical Monitoring



- ☐ Count XXI: Punitive Damages
- ☐ Count XXII: Other [specify below]

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13. As to Philips RS North America Holding Corporation, Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

- ☐ Count I: Negligence
- ☐ Count II: Strict Liability: Design Defect
- ☐ Count III: Negligent Design
- ☐ Count IV: Strict Liability: Failure to Warn
- ☐ Count V: Negligent Failure to Warn
- ☐ Count VI: Negligent Recall
- ☐ Count VII: Battery
- ☐ Count VIII: Strict Liability: Manufacturing Defect
- ☐ Count IX: Negligent Manufacturing
- ☐ Count X: Breach of Express Warranty
- ☐ Count XI: Breach of the Implied Warranty of Merchantability
- ☐ Count XII: Breach of the Implied Warranty of Usability
- ☐ Count XIII: Fraud
- ☐ Count XIV: Negligent Misrepresentation
- ☐ Count XV: Negligence Per Se

- ☐ Count XVI: Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
- ☐ Count XVII: Unjust Enrichment
- ☐ Count XVIII: Loss of Consortium
- ☐ Count XIX: Survivorship and Wrongful Death
- ☐ Count XX: Medical Monitoring
- ☐ Count XXI: Punitive Damages
- ☐ Count XXII: Other [specify below]

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14. As to Polymer Technologies, Inc., Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

- ☐ Count I: Negligence
- ☐ Count II: Strict Liability: Design Defect
- ☐ Count III: Negligent Design
- ☐ Count IV: Strict Liability: Failure to Warn
- ☐ Count V: Negligent Failure to Warn
- ☐ Count VIII: Strict Liability: Manufacturing Defect
- ☐ Count IX: Negligent Manufacturing
- ☐ Count XIII: Fraud
- ☐ Count XIV: Negligent Misrepresentation
- ☐ Count XVII: Unjust Enrichment

- ☐ Count XVIII: Loss of Consortium
- ☐ Count XIX: Survivorship and Wrongful Death
- ☐ Count XX: Medical Monitoring
- ☐ Count XXI: Punitive Damages
- ☐ Count XXII: Other [specify below]

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15. As to Polymer Molded Products LLC, Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

- ☐ Count I: Negligence
- ☐ Count II: Strict Liability: Design Defect
- ☐ Count III: Negligent Design
- ☐ Count IV: Strict Liability: Failure to Warn
- ☐ Count V: Negligent Failure to Warn
- ☐ Count VIII: Strict Liability: Manufacturing Defect
- ☐ Count IX: Negligent Manufacturing
- ☐ Count XIII: Fraud
- ☐ Count XIV: Negligent Misrepresentation
- ☐ Count XVII: Unjust Enrichment
- ☐ Count XVIII: Loss of Consortium
- ☐ Count XIX: Survivorship and Wrongful Death
- ☐ Count XX: Medical Monitoring

☐ Count XXI: Punitive Damages

☐ Count XXII: Other [specify below]

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16. If additional claims against the Defendants identified in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial are alleged above, the additional facts, if any, supporting these allegations must be pleaded. Plaintiff(s) assert(s) the following additional factual allegations against the Defendants identified in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial:

N/A

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17. Plaintiff(s) contend(s) that additional parties may be liable or responsible for Plaintiff(s)' damages alleged herein. Such additional parties, who will be hereafter referred to as Defendants, are as follows (must name each Defendant and its citizenship):

Defendants John Does 1-20-upon information and belief, Defendants John Does 1-20 are entities or persons who are liable to Plaintiff but who have not yet been identified despite reasonable due diligence on the part of Plaintiff. Upon information and belief, Defendants John Does 1-20 (fictitious names) are

entities or persons who are liable to Plaintiff, but who have not yet been identified despite reasonable due diligence on the part of Plaintiff.

26. Upon information and belief, Defendants JOHN DOES 1-20 research, develop, design, manufacture, sell, distribute, and market BiPAP and CPAP devices and ventilators, including the recalled devices and subject devices.

27. Upon information and belief, Defendants JOHN DOES 1-20, researched, developed, designed, manufactured, sold, distributed, and promoted the recalled devices, including the subject devices.

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18. Plaintiff(s) assert(s) the following additional claims and factual allegations against other Defendants named in Paragraph 16 above:

N/A

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WHEREFORE, Plaintiff(s) pray(s) for relief and judgment against Defendants and all such further relief that this Court deems equitable and just as set forth in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial and any additional relief to which Plaintiff(s) may be entitled.

Date: Dec 20 2022

**/s/ Jennifer S. Koockogey-Lajoie**

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